

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
13 February 2003 (13.02.2003)

PCT

(10) International Publication Number
WO 03/011396 A1

(51) International Patent Classification⁷: **A61P 25/00**,
25/18, 25/22, 25/24, 15/00, 15/10, A61K 31/496, 31/506,
C07D 403/12, 209/08

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(21) International Application Number: PCT/US02/24260

(22) International Filing Date: 30 July 2002 (30.07.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/308,644 30 July 2001 (30.07.2001) US
60/365,005 13 March 2002 (13.03.2002) US
60/371,381 9 April 2002 (09.04.2002) US

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(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: TETRAHYDROINDOLONE AND PURINE DERIVATIVES LINKED TO ARYLPIPERAZINES

(57) Abstract: Pharmaceutical composite compositions comprising tetrahydroindolones linked to arylpiperazines and derivatives thereof are disclosed. Specifically, composite compositions useful in treating anti-psychotic disorders are disclosed. The composite compositions disclosed herein can effectively ameliorate symptoms and treat psychotic disorders without causing a decrease in cognitive function. Generally, the composite compounds consist of two moieties, moiety A and B in which a tetrahydroindolone comprises a moiety A linked through a linker L to a moiety B, where B is an arylpiperazinyl moiety. The composite compound provides anti-psychotic activity by interaction with GABA, serotonin and dopamine receptors. The composite molecules with the combined activities will provide treatment of psychiatric and neurological diseases without cognitive impairment.



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TETRAHYDROINDOLONE AND PURINE DERIVATIVES LINKED TO ARYLPYPERAZINES**BACKGROUND OF THE INVENTION****RELATED APPLICATIONS**

[0001] This application claims priority to U.S. provisional application serial number 60/371,381 filed April 9, 2002; U.S. provisional application serial number 60/365,005 filed March 13, 2002; U.S. provisional application serial number 60/308,644 filed July 30, 2001.

FIELD OF THE INVENTION:

[0002] The present invention is directed to compositions comprising tetrahydroindolone linked to arylpiperazines for the treatment of psychosis and psychosis related diseases including cognitive impairment.

BACKGROUND:

[0003] Psychiatric and neurologic disorders are among the most severe and chronic diseases and conditions. These disorders are also extremely difficult to treat effectively because of the multiplicity of the symptoms and etiologies. Although many alternative and combination drug therapies have been attempted and proposed, an effective treatment remains elusive. The current therapies have focused on either high selectivity for one pharmacological effect or broad non-selectivity to attempt multiple symptom relief. Drug therapies that have focused on high pharmacological selectivity have limited benefit for disorders with multiple causes and can worsen some symptoms. For example, selective antagonism of dopaminergic receptors for schizophrenia results in a worsening of negative symptoms and tardive dyskinesia. In contrast to selective drug therapy, drug therapies having broad non-selectivity can provide relief for more symptoms, yet they have more side effects. For instance, current antipsychotic drugs have adrenergic, cholinergic, and histaminergic receptor antagonist activity which are associated with deterioration of cognitive function and other side-effects such as orthostatic hypotension, dry mouth, blurred vision, constipation, and motor impairment. Regardless of drug therapy selectivity, cognitive decline is one main symptom that is not adequately treated in both current psychiatric and neurological therapies.

[0004] Therefore, there is a particular need for the development of compounds that have improved activity in treating psychiatric disorders having symptoms including reduced cognition and emotional control such as schizophrenia, schizoaffective disorders, anxiety and depression with agitation without the side-effects of the existing therapies. There is a further need for compounds that provide treatment or relief from symptoms associated with neurological diseases such as, Alzheimer's disease, movement disorders, (such as Huntington's disease and Parkinson's disease), stroke pain and other neurodegenerative disorders, which can be genetic, spontaneous or iatrogenic. Additionally, there is a need for compounds that are neuroprotective, stimulate neuronal function, neuronal regeneration, neurogenesis and have fewer side effects.

INVENTION SUMMARY

[0005] Generally, the present invention relates to novel composite compounds having the general schematic structure, A}-L-{B, where A is a bicyclic ring structure such as tetrahydroindolone or a purine derivative, L is a hydrocarbonyl chain, and B is an arylpiperazine or arylpiperazine derivative that are useful as pharmaceutical compositions for treating a wide range of psychiatric and neurological disorders.

[0006] Furthermore, these psychiatric and neurological disorders can be effectively treated without causing a deterioration of cognitive function or other major side effects. More particularly, the composite compounds of the present invention comprise tetrahydroindolones or purine derivatives linked to arylpiperazine structures.

[0007] Moreover, the composite compounds of the present invention can have other beneficial effects such as, but not limited to neuroprotective and neuroregenerative properties.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0008] In accordance with the present invention, and as used herein, the following terms, when appearing alone or as part of a moiety including other atoms or groups, are defined with the following meanings, unless explicitly stated otherwise. In addition, all groups described herein can be optionally substituted unless such substitution is excluded. The term "alkyl," as used herein at all occurrences, refers to saturated aliphatic groups including straight-chain, branched-chain, and cyclic groups, all of which can be optionally substituted. Preferred alkyl groups contain 1 to 10 carbon atoms. Suitable alkyl groups include methyl, ethyl, and the like, and can be

optionally substituted. The term "heteroalkyl" as used herein at all occurrences refers to carbon-containing straight-chained, branch-chained and cyclic groups, all of which can be optionally substituted, containing at least one O, N or S heteroatoms. The term "heteroalkenyl" as used herein at all occurrences refers to unsaturated groups which contain at least one carbon-carbon double bond and includes straight-chained, branch-chained and cyclic groups, all of which can be optionally substituted, containing at least one O, N or S heteroatoms. The term "alkenyl," as used herein at all occurrences, refers to unsaturated groups which contain at least one carbon-carbon double bond and includes straight-chain, branched-chain, and cyclic groups, all of which can be optionally substituted. Preferable alkenyl groups have 2 to 10 carbon atoms. The term "alkoxy" refers to the ether --O--alkyl , where alkyl is defined as above. The term "aryl" refers to aromatic groups which have at least one ring having a conjugated π -electron system and includes carbocyclic aryl and biaryl, both of which can be optionally substituted. Preferred aryl groups have 6 to 10 carbon atoms. The term "aralkyl" refers to an alkyl group substituted with an aryl group. Suitable aralkyl groups include benzyl and the like; these groups can be optionally substituted. The term "aralkenyl" refers to an alkenyl group substituted with an aryl group. The term "heteroaryl" refers to carbon-containing 5-14 membered cyclic unsaturated radicals containing one, two, three, or four O, N, or S heteroatoms and having 6, 10, or 14 π -electrons delocalized in one or more rings, e.g., pyridine, oxazole, indole, thiazole, isoxazole, pyrazole, pyrrole, each of which can be optionally substituted as discussed above. The term "sulfonyl" refers to the group $\text{--S(O)}_2\text{--}$. The term "alkanoyl" refers to the group --C(O)Rg , where Rg is alkyl. The term "aroyl" refers to the group --C(O)Rg , where Rg is aryl. Similar compound radicals involving a carbonyl group and other groups are defined by analogy. The term "aminocarbonyl" refers to the group --NHC(O)-- . The term "oxycarbonyl" refers to the group --OC(O)-- . The term "heteroaralkyl" refers to an alkyl group substituted with a heteroaryl group. Similarly, the term "heteroaralkenyl" refers to an alkenyl group substituted with a heteroaryl group. Where used herein, the term "lower," in reference to an alkyl or the alkyl portion of another group including alkyl, is defined as a group containing one to ten carbon atoms, more typically one to six carbon atoms. The term "optionally substituted" refers to one or more substituents that are typically lower alkyl, aryl, amino, hydroxy, lower alkoxy, aryloxy, lower alkylamino, arylamino, lower alkylthio, arylthio, or oxo, in some cases, other groups can be included, such as cyano, acetoxy,

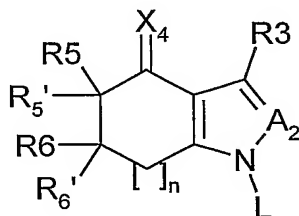
or halo. The term “halo” refers generally to fluoro, chloro, bromo, or iodo; more typically, “halo” refers to chloro.

[0009] In accordance with the present invention, and as used herein, the term “derivative” refers to a compound that is modified or partially substituted with another component. Additionally, the term “derivative” encompasses compounds that can be structurally similar but can have similar or different functions.

[0010] The composite compounds of the present invention have the general schematic structure, A}-L-{B, where A is a bicyclic ring structure such as tetrahydroindolone or a purine derivative, L is a hydrocarbyl chain, and B is an arylpiperazine or arylpiperazine derivative.

I. TETRAHYDROINDOLONE AND PURINE DERIVATIVES

[0011] In one embodiment of the present invention, A is an 8-10 atom bicyclic moiety in which the five-aromatic membered ring has 1 to 2 nitrogen atoms, the bicyclic moiety having the structure of formula (I):



(I)

where:

- (a) formula I is bonded to a hydrocarbyl linker L;
- (b) A₂ is C or N;
- (c) R₃ is hydrogen, alkyl, aralkyl, heteroaralkyl, heteroalkyl, alkenyl, aralkenyl, heteroaralkenyl, heteroalkenyl, aryl, or heteroaryl;
- (d) X₄ is O, S or N-OH;
- (e) R₅ is hydrogen, alkyl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, NH₂, NHQ₁, NQ₁Q₂, OH, OQ₁, or SQ₁, where Q₁ and Q₂ are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and

can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q₁ and Q₂ are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y₂, where Y₂ is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxy carbonyl, heteroaryloxy carbonyl, aralkoxy carbonyl, heteroaralkoxy carbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S;

(f) R_{5'} is hydrogen unless R₅ is alkyl, in which case R_{5'} is hydrogen or the same alkyl as R₅;

(g) R₅ and R_{5'} can be taken together as a double bond to C₅ and can be O, S, NQ₃, or C which can be substituted with one or two groups R₅, where Q₃ is alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, hydroxy, alkoxy, aryloxy, or heteroaryloxy in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S;

(h) R₆ is hydrogen, alkyl, aryl, heteroaryl;

(i) R_{6'} is hydrogen unless R₆ is alkyl, in which case R_{6'} is hydrogen or the same alkyl as R₆;

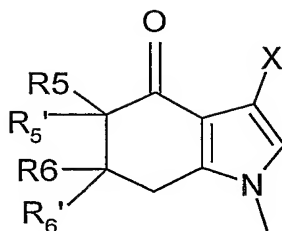
(j) n is 0 to 2.

[0012] As shown in Formula (I), the moiety A has a five, six, or seven-membered saturated ring fused to a five-membered aromatic ring. The five-membered aromatic ring can have one or two nitrogen atoms as indicated, but the five-membered aromatic ring always has a nitrogen atom at the 1-position. Typically, the five-membered aromatic ring has one nitrogen atom as in tetrahydroindolone. This nitrogen atom at the 1-position is covalently bonded to the linker L.

[0013] Typically A is a tetrahydroindolone moiety in which A₂ is carbon and n is 1. The tetrahydroindolone moiety can be variously substituted.

[0014] In yet another embodiment of the present invention, A is a tetrahydroindolone moiety.

[0015] One example of a tetrahydroindolone moiety for the moiety A is a tetrahydroindolone moiety of Formula (II), below, in which:



(II)

(1) Where X is H or $\text{CH}_2\text{N}(\text{CH}_3)_2$;

(2) R_5 is hydrogen, alkyl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, NH_2 , NHW_1 , NQ_1Q_2 , OH, OQ_1 , or SQ_1 , where Q_1 and Q_2 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, or heteroaroyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and where W_1 is alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, or heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S;

(3) $\text{R}_{5'}$ is hydrogen;

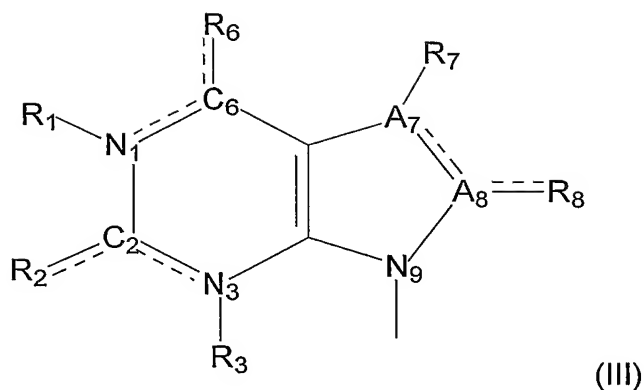
(4) R_6 is hydrogen, alkyl, aryl, heteroaryl;

(5) $\text{R}_{6'}$ is hydrogen.

[0016] In one particularly preferred embodiment, R_5 , $\text{R}_{5'}$, R_6 , and $\text{R}_{6'}$ are all hydrogen. In this particularly preferred embodiment, the moiety A is an unsubstituted tetrahydroindolone moiety.

[0017] In addition to these examples of moieties suitable as moiety A, other moieties can serve as moiety A, including moieties with five or seven-membered saturated rings, and two nitrogen atoms in the five-membered aromatic ring or moieties with substituents at R_3 .

[0018] In another embodiment of the present invention, A is a substituted or unsubstituted 9-atom bicyclic moiety in which the five-membered ring has 1 to 3 nitrogen atoms, the bicyclic moiety having the structure of formula (III)



where:

(a) if the bond between N₁ and C₆ is a single bond, then the bond between C₆ and R₆ is a double bond, R₆ is O or S, and R₁ is hydrogen, alkyl, aralkyl, cycloalkyl, or heteroaralkyl;

(b) if the bond between N₁ and C₆ is a double bond, then the bond between C₆ and R₆ is a single bond, R₁ is not present, and R₆ is hydrogen, halo, amino, OQ₁, SQ₁, NHNH₂, NHOQ₁, NQ₁Q₂, or NHQ₁, where Q₁ and Q₂ are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q₁ and Q₂ are present together and are alkyl, they can be taken together to form a 5 or 6 member ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y₂, where Y₂ is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S;

(c) if the bond between C₂ and N₃ is a single bond, then the bond between C₂ and R₂ is a double bond, R₂ is O or S, and R₃ is hydrogen or alkyl;

(d) if the bond between C₂ and N₃ is a double bond, then the bond between C₂ and R₂ is a single bond, R₃ is not present, and R₂ is hydrogen, alkyl, aralkyl, cycloalkyl, heteroaralkyl, halo, amino, OQ₁, SQ₁, NHNH₂, NHOQ₁, NQ₁Q₂, or NHQ₁, where Q₁ and Q₂ are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl,

aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q₁ and Q₂ are present together and are alkyl, they can be taken together to form a 5 or 6 member ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y₂, where Y₂ is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S;

(e) A₇ and A₈ are C or N;

(i) if A₇ and A₈ are both C and the bond between A₇ and A₈ is a single bond, then the bond between A₈ and R₈ is two single bonds to two hydrogen atoms or is a double bond in which R₈ is O or S and R₇ is two hydrogen atoms;

(ii) if A₇ and A₈ are both C and the bond between A₇ and A₈ is a double bond, then R₇ is hydrogen, the bond between A₈ and R₈ is a single bond and R₈ is hydrogen, halo, alkyl, alkenyl, aryl, aralkyl, aralkenyl, heteroaryl, heteroaralkyl, or heteroaralkenyl;

(iii) if A₇ and A₈ are both N, then the bond between A₇ and A₈ is a double bond and R₇ and R₈ are not present;

(iv) if A₇ is C and A₈ is N, then the bond between A₇ and A₈ is a double bond, R₇ is hydrogen, and R₈ is not present;

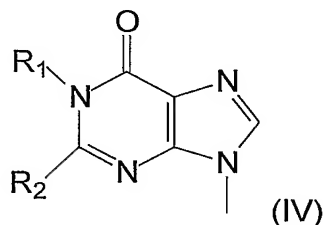
(v) if A₇ is N, A₈ is C, and the bond between A₇ and A₈ is a double bond, then R₇ is not present, the bond between A₈ and R₈ is a single bond, and R₈ is hydrogen, halo, alkyl, alkenyl, aryl, aralkyl, aralkenyl, heteroaryl, heteroaralkyl, or heteroaralkenyl;

(vi) if A₇ is N, A₈ is C, and the bond between A₇ and A₈ is a single bond, then R₇ is hydrogen, alkyl, aryl, aralkyl, heteroaryl, or heteroaralkyl, the bond between A₈ and R₈ is a double bond, and R₈ is O or S;

(f) N₉ is bonded to linker L.

[0019] Particularly preferred purine moieties for the moiety A are described below.

[0020] One example of a purine moiety for the moiety A is a purine moiety of Formula (IV), below, in which:



R_1 is selected from the group consisting of hydrogen, alkyl, aralkyl, cycloalkyl, heteroaralkyl, and

R_2 is selected from the group consisting of hydrogen, alkyl, aralkyl, cycloalkyl, heteroaralkyl, halo, amino, OQ_1 , SQ_1 , $NHNH_2$, $NHOQ_1$, NQ_1Q_2 , or NHQ_1 , where Q_1 and Q_2 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q_1 and Q_2 are present together and are alkyl, they can be taken together to form a 5 or 6 member ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y_2 , where Y_2 is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxy carbonyl, heteroaryloxy carbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S.

[0021] The purine moiety of formula (IV) is a hypoxanthine or a guanine derivative.

[0022] In one particularly preferred embodiment, R_1 is hydrogen and R_2 is hydrogen.

[0023] In another preferred embodiment, R_1 is hydrogen and R_2 is amino.

[0024] In one particularly preferred embodiment, R_1 is butyl and R_2 is hydrogen.

[0025] In another preferred embodiment, R_1 is benzyl and R_2 is hydrogen.

[0026] In another preferred embodiment, R_1 is dimethylaminoethyl and R_2 is hydrogen.

[0027] In another preferred embodiment, R_1 is cyclopentyl and R_2 is hydrogen.

[0028] In another preferred embodiment, R_1 is cyclohexylmethyl and R_2 is hydrogen.

[0029] In another preferred embodiment, R_1 is cyclopropylmethyl and R_2 is hydrogen.

[0030] In another preferred embodiment, R_1 is hydrogen and R_2 is phenyl.

[0031] In another preferred embodiment, R_1 is hydrogen and R_2 is trifluoromethyl.

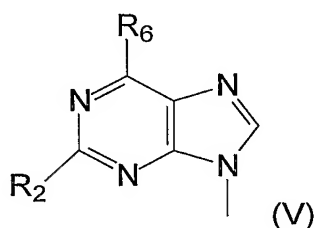
[0032] In another preferred embodiment, R_1 is hydrogen and R_2 is butyl.

[0033] In another preferred embodiment, R_1 is butyl and R_2 is butyl.

[0034] In another preferred embodiment, R_1 is hydrogen and R_2 is methyl.

[0035] In another preferred embodiment, R_1 is hydrogen and R_2 is phenylamino.

[0036] Another example of a purine moiety according to the present invention is the purine moiety of Formula (V), below, in which:



R_2 is selected from the group consisting of hydrogen, halo, amino, OQ_3 , SQ_3 , $NHNH_2$, $NHOQ_3$, NQ_3Q_4 , or NHQ_3 , where Q_3 and Q_4 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, and heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q_3 and Q_4 are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which can contain one other heteroatom which can be N, O, or S, of which the N can be further substituted with Y_3 , where Y_3 is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylamino carbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S; and

R_6 is selected from the group consisting of hydrogen, halo, amino, OQ_5 , SQ_5 , $NHNH_2$, $NHOQ_5$, NQ_5Q_6 , or NHQ_5 , where Q_5 and Q_6 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, and

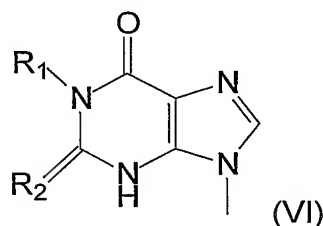
heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q₅ and Q₆ are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which can contain one other heteroatom which can be N, O, or S, of which the N can be further substituted with Y₂, where Y₂ is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S.

[0037] In one preferred example of this embodiment, R₂ is hydrogen and R₆ is -NH₂ or -N(CH₃)₂.

[0038] In another preferred example of this embodiment, R₂ is hydrogen and R₆ is Cl.

[0039] In yet another preferred example of this embodiment, R₂ is -NH₂ and R₆ is Cl.

[0040] Another example of a purine moiety according to the present invention is the purine moiety of Formula (VI), below, in which:



R₁ is hydrogen, alkyl, aralkyl, cycloalkyl, or heteroaralkyl.

R₂ is O or S.

[0041] Preferably, R₁ is hydrogen.

[0042] The purine moiety of Formula (VI) is a xanthine or thioxanthine moiety.

[0043] In addition to these examples of moieties suitable as moiety A, other moieties can serve as moiety A, including moieties with one or three nitrogen atoms in the five-membered ring or moieties with substituents at R₈.

[0044] Typically, for purine derivatives according to the present invention, A is a hypoxanthine moiety.

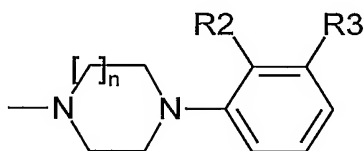
II. HYDROCARBYL LINKER

[0045] In one embodiment, the linker L is a hydrocarbyl moiety with the structure $-(CH_2)_m-$ wherein m is an integer from 1 to 6. A preferred linker has m equal to 2, 3 or 4.

[0046] In another embodiment, linker L is a phenyl or a benzyl linked to a hydrocarbyl chain by group Y where group Y is located on the meta or para positions of the aromatic ring. Group Y can be nothing such that the hydrocarbyl chain is directly linked to the phenyl group. Group Y can also be an ether, thioether, carbonyl, thiocarbonyl, carboxamido, aminocarbonyl, amino, oxycarbonylamino, aminocarbonyloxy, aminocarbonylamino, oxythiocarbonylamino, aminothiocabonyloxy, aminothiocabonylamino, aminosulfonyl, or sulfonamido group.

III. ARYLPIPERAZINE AND DERIVATIVES

[0047] B is an arylpiperazine or derivative having the structure of formula (VII):

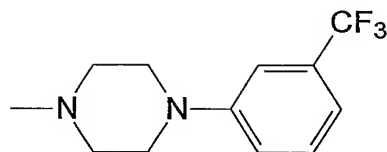


(VII)

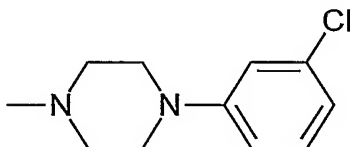
where:

- (a) R2 is hydrogen, alkyl, hydroxy, halo, alkoxy, cyano, methylthio;
- (b) R3 is hydrogen, alkyl, hydroxy, methoxy, halo, alkoxy, trifluoromethyl, nitro, amino, aminocarbonyl, aminosulfonyl;
- (c) R2 and R3 can be taken together to form a 5 or 6 member aromatic or non-aromatic ring, which can contain from 0 to 3 heteroatoms selected from the group of N, O, or S.
- (d) n equals 1 or 2

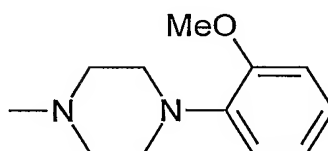
[0048] In one embodiment, B is a *m*-trifluoromethylphenylpiperazinyl moiety:



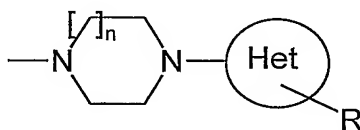
[0049] In another embodiment, B is a *m*-chlorophenylpiperazinyl moiety:



[0050] In yet another embodiment, B is an *o*-methoxyphenylpiperazinyl moiety:



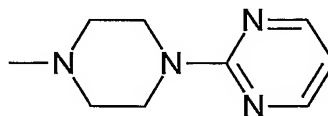
[0051] In another embodiment, B is a piperazine ring or derivative linked to a 6-member heterocyclic ring containing 1 to 3 N, having the structural formula (VIII):



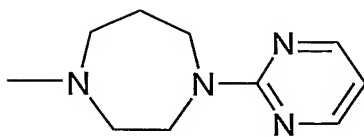
(VIII)

Wherein $n = 1$ or 2 and the 6-member heterocyclic ring (Het) can be 2-pyridyl, 4-pyridyl, 2-pyrimidyl, 4-pyrimidyl, 2-pyrazinyl, or 2-triazinyl. The heterocyclic ring can also be substituted where R can be halo, alkyl, cyano, trifluoromethyl, alkoxy, amino, alkylamino, or dialkylamino.

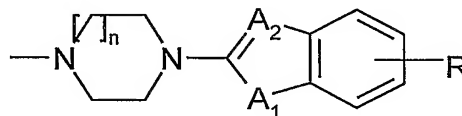
[0052] In one embodiment, B is a 2-pyrimidylpiperazinyl moiety:



[0053] In another embodiment, B is a 1-pyrimidin-2-yl-[1,4]diazepane moiety:



[0054] In yet another embodiment, B is a piperazine ring or derivative linked to a bicyclic moiety having the structural formula (IX):



(IX)

where:

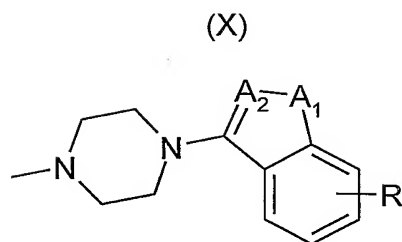
(a) A₁ is N, O, or S, and when it is N, it can be further substituted with Z, which is alkyl, aralkyl, heteroaralkyl, or heteroalkyl.

(b) A₂ is C or N;

(c) and n is 1 or 2

(d) R is hydrogen, alkyl, NH₂, NHQ₁, NQ₁Q₂, OH, OQ₁, SQ₁, halo, nitro, cyano, or trifluoromethyl where Q₁ and Q₂ are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q₁ and Q₂ are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y₂, where Y₂ is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S.

In another embodiment, B is piperazine ring or derivative linked to a bicyclic moiety having the structure (X) below:



where:

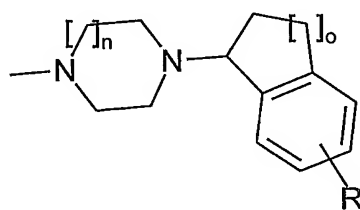
(a) A_1 is N, O, or S, and when it is N, it can be further substituted with Z, which in alkyl, aralkyl, heteroaralkyl, or heteroalkyl.

(b) A_2 is C or N;

(c) and n is 1 or 2

(d) R is hydrogen, alkyl, NH_2 , NHQ_1 , NQ_1Q_2 , OH, OQ_1 , SQ_1 , halo, nitro, cyano, or trifluoromethyl where Q_1 and Q_2 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q_1 and Q_2 are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which may contain 1 other heteroatom which can be N, O, or S, of which the N may be further substituted with Y_2 , where Y_2 is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylaminocarbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S.

[0055] In another embodiment, B is a piperazine ring or derivative linked to a bicyclic moiety having the structural formula (X):

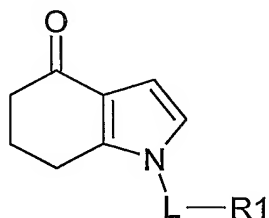


(XI)

where:

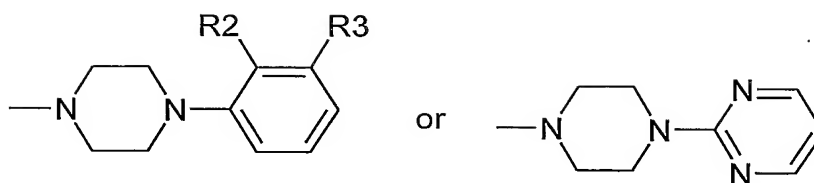
- (a) o is 1 to 3;
- (b) n is 1 or 2; and
- (c) R is hydrogen, alkyl, NH_2 , NHQ_1 , NQ_1Q_2 , OH, OQ_1 , SQ_1 , nitro, cyano, trifluoromethyl, or halo where Q_1 and Q_2 are alkyl, aralkyl, heteroaralkyl, aryl, heteroaryl, alkanoyl, aroyl, aralkanoyl, heteroaralkanoyl, heteroaroyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, or heteroaralkylsulfonyl in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S, and when Q_1 and Q_2 are present together and are alkyl, they can be taken together to form a 5- or 6-membered ring which can contain 1 other heteroatom which can be N, O, or S, of which the N can be further substituted with Y_2 , where Y_2 is alkyl, aryl, heteroaryl, aralkyl, heteroaralkyl, alkanoyl, aroyl, heteroaroyl, aralkanoyl, heteroaralkanoyl, alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, aralkylsulfonyl, heteroaralkylsulfonyl, alkoxycarbonyl, aryloxycarbonyl, heteroaryloxycarbonyl, aralkoxycarbonyl, heteroaralkoxycarbonyl, alkylaminocarbonyl, arylaminocarbonyl, heteroarylamino carbonyl, aralkylaminocarbonyl, or heteroaralkylaminocarbonyl, in which the alkyl portions can be cyclic and can contain from 1 to 3 heteroatoms which can be N, O, or S.

[0056] Generally, any moiety A can be combined with any linker L and any moiety B to produce a composite compound according to the present invention. However, in one embodiment the composite compounds of the present invention include, but are not limited to, the following structure:



(XII)

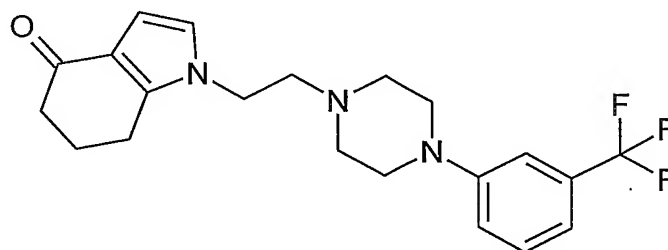
- (a) wherein L is $-(\text{CH}_2)_m-$ wherein m is an integer from 1 to 6;
- (b) R1 is:



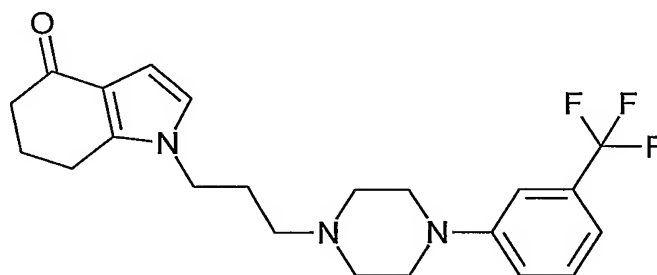
and R2 and R3 are the same or independently hydrogen, alkyl, hydroxy, methoxy, halo, alkoxy, trifluoromethyl, nitro, amino, aminocarbonyl, or aminosulfonyl.

[0057] More specifically, the composite compounds of the present invention include, but are not limited to:

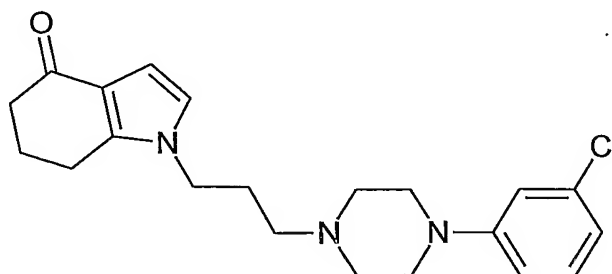
- (1) 1-{2-[4-(3-Trifluoromethylphenyl)piperazine-1-yl]ethyl}-1,5,6,7-tetrahydroindol-4-one (NEO-359)



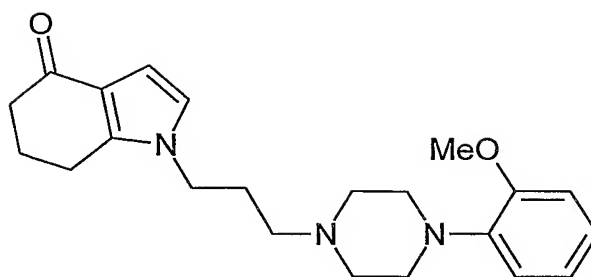
- (2) 1-{3-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-356)



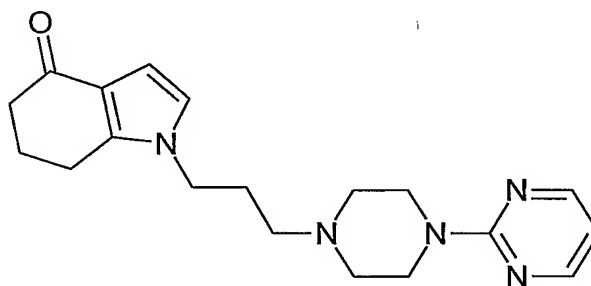
- (3) 1-{3-[4-(3-Chlorophenyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-363)



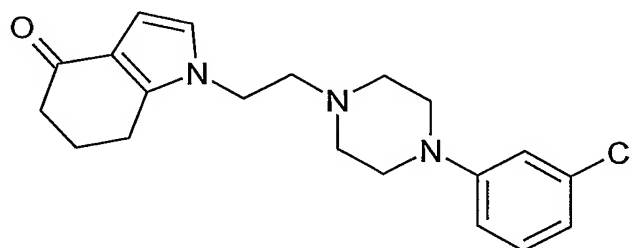
- (4) 1-{3-[4-(2-Methoxyphenyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-370)



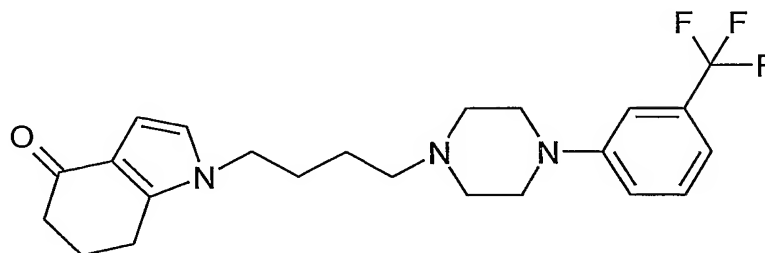
(5) 1-{3-[4-(2-Pyrimidyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-381)



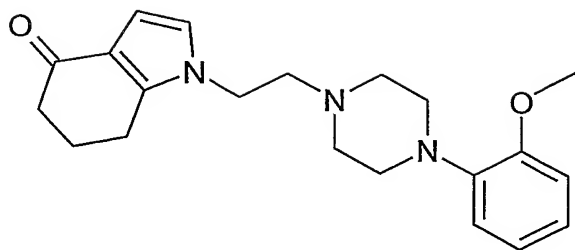
(6) 1-{2-[4-(3-Chloro-phenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydroindol-4-one (NEO-376)



(7) 1-{4-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]-butyl}-1,5,6,7-tetrahydroindol-4-one (NEO-392)

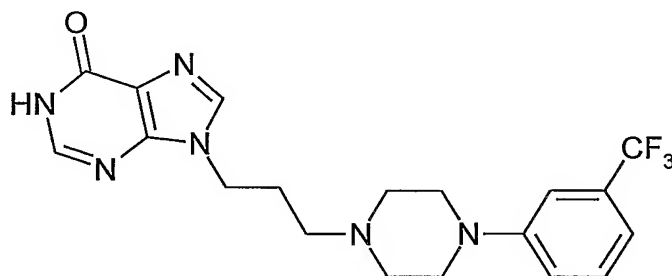


(8) 1-{2-[4-(2-Methoxyphenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydroindol-4-one (NEO-377)

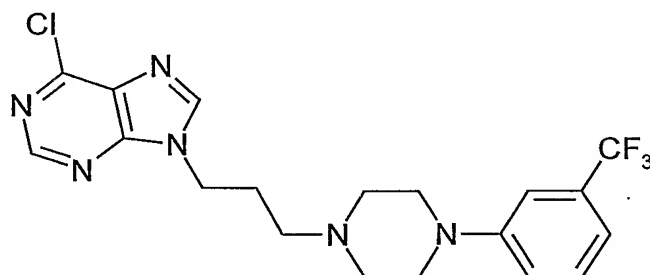


[0058] Other examples of the composite compounds of the present invention include, but are not limited to:

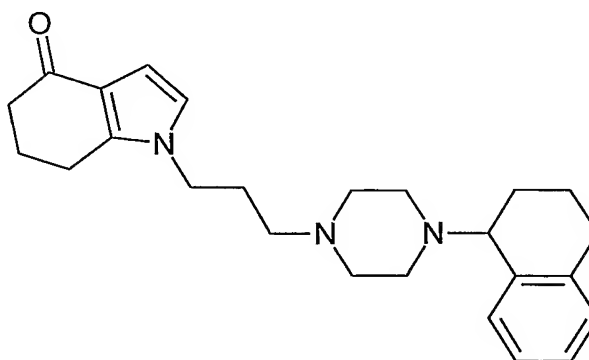
(9) 9-{3-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]propyl}-1,9-dihydropurin-6-one



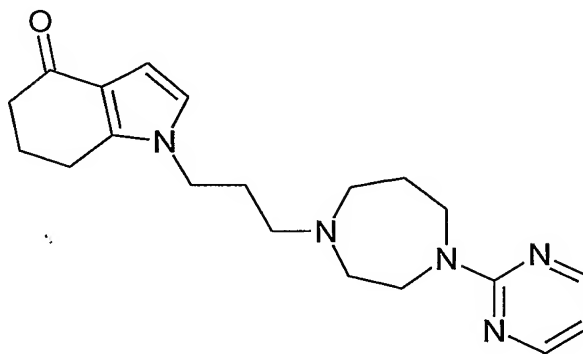
(10) 6-Chloro-9-{3-[4-(3-trifluoromethylphenyl)piperazin-1-yl]propyl}-9H-purine



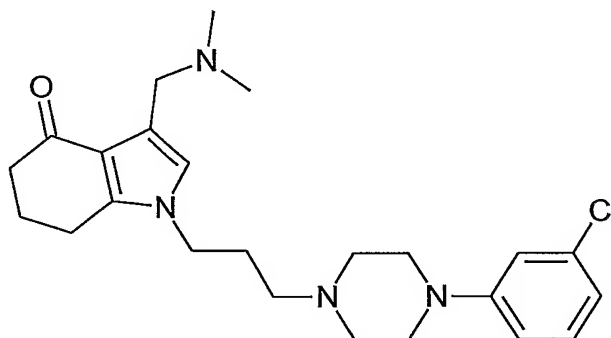
(11) 1-{3-[4-(1,2,3,4-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-propyl}-1,5,6,7-tetrahydro-indol-4-one



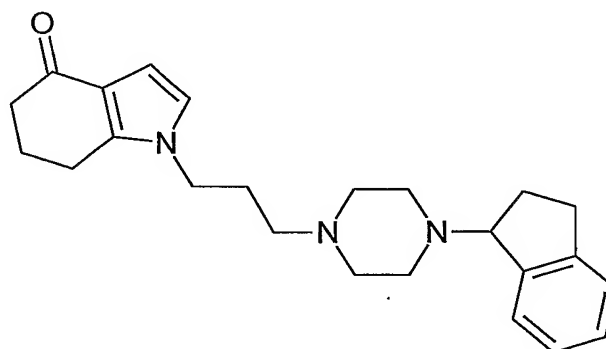
(12) 1-[3-(4-Pyrimidin-2-yl-[1,4]diazepan-1-yl)-propyl]-1,5,6,7-tetrahydro-indol-4-one



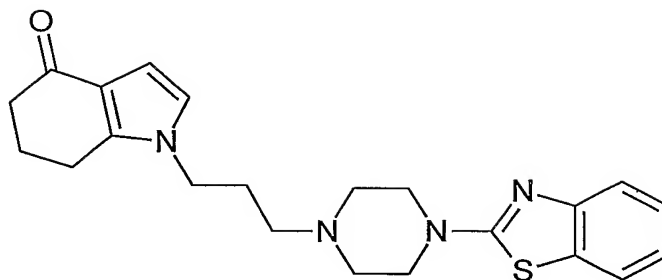
(13) 1-{3-[4-(3-Chloro-phenyl)-piperazin-1-yl]-propyl}-3-dimethylaminomethyl-1,5,6,7-tetrahydro-indol-4-one



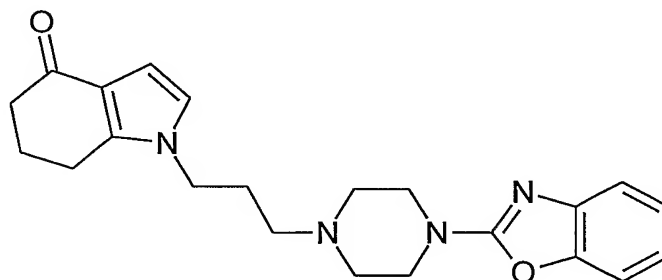
(14) 1-{3-[4-(1,2,3,4-Tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-propyl}-1,5,6,7-tetrahydro-indol-4-one



15) 1-[3-(4-Benzothiazol-2-ylpiperazin-1-yl)propyl]-1,5,6,7-tetrahydroindol-4-one



(16) 1-[3-(4-Benzoxazol-2-yl)piperazin-1-yl]propyl-1,5,6,7-tetrahydroindol-4-one



[0059] According to the present invention, a tetrahydroindolone or purine derivative linked to moiety B is referred to as a “composite compound.” According to the present invention, moiety B has a biological, physiological, or pharmacological function, but it is not required that moiety B of the composite compound have a biological, physiological, or pharmacological function. The moiety B can serve as a carrier to improve bioavailability or to optimize the physical characteristics of the molecule without having a separate biological, physiological function, or pharmacological function.

[0060] Typically, the composite compound has a logP of from about 1 to about 4 to enhance bioavailability and central nervous system (CNS) penetration. Using this guideline, one of ordinary skill in the art can choose the appropriate moieties B for a particular moiety A in order to ensure the bioavailability and CNS penetration of the composite compound of present invention. For example, if a highly hydrophobic moiety A is chosen, with particularly hydrophobic substituents on the tetrahydroindolone or purine moiety, then a more hydrophilic moiety B can be used.

[0061] A number of composite compounds according to the present invention are optically active, owing to the presence of chiral carbons or other centers of asymmetry. In cases where composite compounds of the present invention are

optically active, all of the possible enantiomers or diastereoisomers are included unless otherwise indicated despite possible differences in activity.

[0062] In general, composite compounds that are within the scope of the present invention also include salts and prodrug esters of these derivatives. It is well known that organic compounds, including substituted tetrahydroindolones, purines, arylpiperazines and other components of these composite compounds, have multiple groups that can accept or donate protons, depending upon the pH of the solution in which they are present. These groups include carboxyl groups, hydroxyl groups, amino groups, sulfonic acid groups, and other groups known to be involved in acid-base reactions. The recitation of a composite compound includes such salt forms as occur at physiological pH or at the pH of a pharmaceutical composition unless specifically excluded.

[0063] Similarly, prodrug esters can be formed by reaction of either a carboxyl or a hydroxyl group on the composite compound with either an acid or an alcohol to form an ester. Typically, the acid or alcohol includes a lower alkyl group such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, and tertiary butyl. These groups can be substituted with substituents such as hydroxy, halo, or other substituents. Such prodrugs are well known in the art and need not be described further here. The prodrug is converted into the active compound by hydrolysis of the ester linkage, typically by intracellular enzymes. Other suitable groups that can be used to form prodrug esters are well known in the art.

[0064] Another aspect of the present invention is methods of use of the composite compounds described above. The therapeutic effects of the composite compounds of the present invention are achieved by the effects mediated by multiple receptors. That is, by providing multiple pharmacophores having different receptor activities, the present invention is able to achieve therapeutic benefits without adverse side effects. For example, composite compounds with the arylpiperazine component have serotonin (5-HT) and dopamine activity. In particular, the arylpiperazine component of the composite molecules will have 5-HT₁, and specifically 5TH_{1A} receptor agonist activity, with or without antagonism at 5-HT₂ and D₂, D₃, and D₄ dopamine receptors. Accordingly, composite compounds with all of these activities will be useful in treating schizophrenia, schizoaffective disorders, depression with agitation, and Huntington's disease. These compounds will have fewer side-effects compared to current antipsychotics due to a lack of adrenergic, cholinergic, and histaminergic receptor

antagonist activity. Other composite compounds from this series have 5-HT_{1A} receptor agonist activity and with less selective activity at other receptors. These will be useful as antidepressants, anxiolytics, neuroprotectants and antiemetics.

[0065] Additionally, the tetrahydroindolone or purine derivative component of the composite compound is a pharmacophore having GABA activity. As those skilled in the art will appreciate, GABA receptors are highly localized in the hippocampal region of the brain which is associated with memory. GABA receptors may also be involved in psychosis. Accordingly, it is assumed that pharmacophores, such as tetrahydroindolone or purine derivatives, having GABA activity will improve cognitive function and enhance the therapeutic properties of the arylpiperazine moiety. GABA activity alone will be useful for improving cognition in patients.

[0066] An additional aspect of a method of use of composite compounds according to the present invention is a method of stimulating neuronal function such as improved cognition, involving neurogenesis, neuronal regeneration or axo-dendritic complexity in the central and peripheral nervous systems comprising the step of administering an effective amount of a composite compound according to the present invention to the mammal.

[0067] An additional aspect of a method of use of composite compounds according to the present invention is a method of stimulating neuronal function involving mechanism associated with neuroprotection in the central or peripheral nervous system of a mammal comprising the step of administering an effective amount of the described composite molecules according to the present invention to the mammal.

II. METHODS OF SYNTHESIS OF COMPOSITE MOLECULES ACCORDING TO THE PRESENT INVENTION

[0068] Methods for synthesis of purine and tetrahydroindolone derivatives according to the present invention are modified from those described, for example, in U.S. Patent No. 5,091,432 to Glasky, incorporated herein by this reference. Generally, the tetrahydroindolone moiety is substituted with a linker which in turn is linked to the moiety B that completes the molecule as described above. This route comprises, in either order, the steps of: (1) synthesizing an appropriately substituted tetrahydroindolone moiety linked to an aliphatic linker in which the linker is terminated with a halogen; (2) reacting the halogen intermediate with the arylpiperazine to produce the final product. A similar route can be used for synthesis of purine

derivatives according to the present invention using an appropriately substituted purine moiety as a starting material.

[0069] The length of the aliphatic linker covalently bound to the tetrahydroindolone or purine moiety can be varied to change the distance between the tetrahydroindolone or purine moiety and the moiety B in the composite compound of the present invention.

[0070] Composite compounds of the present invention incorporating an aryl piperazinyl moiety can be synthesized by a dihalide substitution reaction. Suitable substitution reactions are described, e.g., in M.B. Smith & J. March, "March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure" (5th ed., Wiley-Interscience, New York, 2001).

[0071] Another reaction that can be used to functionalize tetrahydroindolones or purines is the Mitsunobu reaction. The Mitsunobu reaction is a highly versatile method for the introduction of widely varying functionality upon the tetrahydroindolone or purine moiety, because of the wide assortment of primary alcohols that are commercially available for use in this reaction.

[0072] Alternatively, in those embodiments where the composite compounds of the present invention include a purine moiety, the purine ring can be formed in stages, with the attachment of the linker and the moiety B occurring before the closure of the purine ring. This route involves: (1) the formation of aminocyanacetamide; (2) the reaction of aminocyanacetamide with triethyl orthoformate and acetonitrile to form an amidoester derivative of aminocyanacetamide; (3) the formation of a compound having a reactive amino group on a hydrocarbyl moiety, the hydrocarbyl moiety being linked to the moiety B; (4) the reaction of the amidoester with the compound having the reactive amino group on the hydrocarbyl moiety; (5) formation of the six-membered heterocyclic ring of the purine moiety; and (6) hydrolysis of the protecting group, if present, to form the final product.

[0073] Additionally, where the composite compounds of the present invention include a purine moiety, in one alternative, the step of the formation of the six-membered heterocyclic ring of the purine moiety can be performed by methods analogous to the ring closure of Yamazaki (A. Yamazaki et al., "Synthesis of Guanosine and Its Derivatives from 5-Amino-1- β -D-Ribofuranosyl-4-Imidazolecarboxamide I. Ring Closure with Benzoyl Isothiocyanate," J. Org. Chem. 32: 1825-1828 (1967)) or alternatively, by the method of Clausen (B. Alhede *et al.*, "A Simple and Efficient Synthesis of 9-Substituted Guanines. Cyclodesulfurization of 1-Substituted 5-

[(Thiocarbamoyl)amino]imidazole-4-Carboxamides Under Aqueous Basic Conditions,” J. Org. Chem. 56: 2139-2143 (1991)) involving catalysis by a heavy metal salt such as Cu^{2+} , Ag^{+} , or Hg^{2+} in aqueous NaOH, or, alternatively, by S-oxidation with hydrogen peroxide or sodium perborate in aqueous sodium hydroxide.

EXAMPLES

[0074] The following representative methods for synthesizing exemplarity embodiments of the present invention are merely intended as examples. Persons having ordinary skill in the art of medicinal and/or organic chemistry will understand that other starting materials, intermediates, reaction conditions are possible. The present examples represent but one particular method for synthesizing the composite, biologically active molecules of the present invention. Furthermore, it is understood that various salts of these compounds are also easily made and these salts can have biological activity similar or exactly equivalent to the parent compound. Generally, these salts have potassium, sodium or calcium as the cation. However, other cations can be used and are considered within the scope of the present invention.

EXAMPLE 1

Synthesis of 1-{2-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]ethyl}-1,5,6,7-tetrahydroindol-4-one (NEO-359)

[0075] This example demonstrates a method of preparing 1-{2-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]ethyl}-1,5,6,7-tetrahydroindol-4-one by a two step procedure. Generally, the arylpiperazine moieties are prepared first, then the arylpiperazine molecules are reacted with tetrahydroindolones.

[0076] Step 1: Preparation of 1-(2-Chloroethyl)-4-(3-trifluoromethylphenyl)piperazine

[0077] To a 100 mL flask was added 4-(3-trifluoromethylphenyl)piperazine HCl (5035 mg, 18.88 mmol) and 60 mL dichloromethane. 1-Bromo-2-chloroethane (1730 μL , 20.78 mmol, 1.10 eq) was added, then triethylamine (5.25 mL, 37.7 mmol, 2.00 eq). The solution was refluxed for 9 hours, then cooled to 25° C. 100 mL of hexane was then added, and the resulting suspension was vacuum filtered. The filtrate was concentrated in vacuum and purified by column chromatography using dichloromethane as eluant resulting in an oil of 1-(2-Chloroethyl)-4-(3-trifluoromethylphenyl)piperazine.

[0078] Step 2: Preparation of 1-{2-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]ethyl}-1,5,6,7-tetrahydroindol-4-one

[0079] Sodium hydride (60% in oil) (85 mg, 2.1 mmol, 1.8 eq.) was added to a 10 ml pear-shaped flask. The solid was rinsed twice with 2 mL hexane to remove oil, then 3 mL anhydrous N,N-dimethylformamide (DMF) was added. 1,5,6,7-Tetrahydroindol-4-one (186.7 mg, 1.38 mmol, 1.159 eq.) was added slowly, with stirring and hydrogen evolved. The walls of the flask were washed with an additional 1 mL of anhydrous DMF. 1-(2-Chloro-ethyl)-4-(3-trifluoromethyl-phenyl)-piperazine (349.00 mg, 1.19 mmol, 1.000 eq) was added as a solution in 2 mL DMF, and the mixture was stirred under nitrogen at 25 C for 8 hours. The resulting mixture was acidified with 1 N HCl to pH 6, and extracted with dichloromethane. The organic layer was washed four times with 25 mL water, dried over sodium sulfate and concentrated in vacuum to an oil which was purified by column chromatography using 5% methanol in dichloromethane as eluant resulting in the title compound as an oil. The oil was dissolved in 5 mL of 50% dichloromethane in hexanes. A solution of 4N HCl in dioxane (200 μ L) was added and the mixture stirred for 30 minutes followed by vacuum filtration of the suspension. A white powder of the product HCl salt was recovered.

EXAMPLE 2

Synthesis of 1-{3-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-356)

[0080] Step 1: Preparation of 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine

[0081] To a 100 mL flask was added 1-(3-trifluoromethylphenyl)piperazine HCl (5035 mg, 18.88 mmol) and 60 mL dichloromethane. 1-Bromo-3-chloropropane (1730 μ L, 20.78 mmol, 1.10 eq) was added, then triethylamine (5.25 mL, 37.7 mmol, 2.00 eq). The solution was refluxed for 9 hours, then cooled to 25° C. 100 mL of hexane was then added, and the resulting suspension was vacuum filtered. The filtrate was concentrated in vacuum and purified by column chromatography using dichloromethane as eluant resulting in an oil of 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine.

[0082] Step 2: Preparation of 1-{2-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one

[0083] The compound is synthesized by reacting the 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine with 1,5,6,7-Tetrahydroindol-4-one using step 2 of Example 1.

EXAMPLE 3

Synthesis of 1-{3-[4-(3-Chlorophenyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-363)

[0084] Since 1-(3-Chloropropyl)-4-(3-chlorophenyl)piperazine HCl is commercially available, step one was omitted.

[0085] To a solution of 1,5,6,7-Tetrahydroindol-4-one (135 mg, 1.0 mmol) in 5 mL dimethylsulfoxide was added powdered sodium hydroxide (84 mg, 2.1 mmol) and the solution stirred for 15 minutes at 25 C. 1-(3-Chloropropyl)-4-(3-chlorophenyl)piperazine HCl (310 mg, 1.0 mmol) was then added and stirring continued overnight. Upon completion, by TLC, reaction was partitioned between 50 mL each of dichloromethane and water then separated. The water layer was extracted with 50 mL more of dichloromethane and the combined organic layers washed with brine, dried with sodium sulfate, and concentrated in vacuum to dryness. The crude product was purified via flash chromatography eluting with an ethyl acetate and dichloromethane mixture resulting in the title compound as an oil. The oil was dissolved in 5 mL of 50% dichloromethane in hexanes. A solution of 4N HCl in dioxane (200 μ L) was added and the mixture stirred for 30 minutes followed by vacuum filtration of the suspension. A white powder of the product HCl salt was recovered.

EXAMPLE 4

Synthesis of 1-{3-[4-(2-Methoxyphenyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-470)

[0086] Step 1: Preparation of 1-(3-Chloropropyl)-4-(2-methoxyphenyl)piperazine

[0087] The 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine is prepared by the same method as disclosed in step 1 of example 2 employing 1-(2-Methoxyphenyl)piperazine HCl instead.

[0088] Step 2: Preparation of 1-{3-[4-(2-Methoxyphenyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one

[0089] The compound is prepared by the same method as disclosed in step 2 of example 3.

EXAMPLE 5

Synthesis of 1-{3-[4-(2-Pyrimidyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one (NEO-381)

[0090] Step 1: Preparation of 1-(3-Chloropropyl)-4-(2-pyrimidyl)piperazine

[0091] The compound is prepared by the same method as disclosed in step 1 of example 2 employing 1-(2-Pyrimidyl)piperazine.2HCl instead.

[0092] Step 2: Preparation of 1-{3-[4-(2-Pyrimidyl)piperazine-1-yl]propyl}-1,5,6,7-tetrahydroindol-4-one

[0093] The compound is prepared by the same method as disclosed in step 2 of example 3.

EXAMPLE 6

Synthesis of 1-{2-[4-(3-Chloro-phenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydro-indol-4-one (NEO-376)

[0094] Step 1: Preparation of 1-(2-Chloroethyl)-4-(3-chlorophenyl) piperazine

[0095] A mixture of (3-chloro-phenyl)-piperazine hydrochloride (51.5 mmol) and powdered sodium hydroxide (103 mmol) in DMSO (75 mL) was treated with 2-bromo-1-chloroethane (77.2 mmol) and stirred at ambient temperature for 4 h. The reaction was poured into ice cold water (200 mL) and stirred for 0.5 h. A solid mass formed and was separated by decanting the water. The aqueous layer was extracted with dichloromethane (100 mL). The solid mass was dissolved with dichloromethane (100 mL) and the combined organics were dried with sodium sulfate, filtered and the solvent removed under vacuum. Flash chromatography (chloroform:acetone 50:1 to 20:1) yielded an oil (7.95 g) as the titled compound.

[0096] Step 2: 1-{2-[4-(3-Chloro-phenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydro-indol-4-one

[0097] To a solution of 1,5,6,7-tetrahydro-indol-4-one (51.5 mmol) in DMSO (60 mL) was added powdered sodium hydroxide (53.9 mmol) and the mixture was stirred at ambient temperature for 0.5 h. 1-(2-Chloroethyl)-4-(3-chlorophenyl) piperazine (49.0 mmol) was then added as a solution in DMSO (20 mL) and the resulting mixture stirred at ambient temperature for 24 h then heated to approximately 60° C for 2 h, after which time TLC (ethyl acetate:dichloromethane 1:1) showed complete reaction. The reaction was poured into ice cold water (300 mL) and stirred for 0.5 h. A solid mass formed and was separated by decanting the water. The aqueous layer was extracted with dichloromethane (100 mL). The solid mass was dissolved with dichloromethane (100 mL) and the combined organics were dried with sodium sulfate and the solvent removed under vacuum. The resulting sludge was triturated with hexanes (100 mL) for 2 h and the suspension vacuum filtered and washed with

hexanes. The obtained solid was under vacuum resulting in a tan powder (14.57 g) as the titled compound, NEO-376).

EXAMPLE 7

Synthesis of 1-{2-[4-(2-Methoxyphenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydro-indol-4-one (NEO-377)

[0098] Step 1: Preparation of 1-(2-Chloroethyl)-4-(2-methoxyphenyl)piperazine

[0099] A mixture of 1-(2-methoxyphenyl)piperazine HCl (52.5 mmol) and powdered sodium hydroxide (105 mmol) in DMSO (40 mL), was stirred at ambient temperature. After 0.5 h, 1-bromo-2-chloroethane (78.8 mmol) was added to the solution and left to stir for 4 h. The reaction was monitored by TLC (ethyl acetate: dichloromethane 1:4), upon completion, the mixture was poured into 200 mL of ice water and the product was extracted with dichloromethane twice, dried with sodium sulfate, and solvent was removed under vacuum. Flash chromatography (ethyl acetate: dichloromethane, 1:5) yielded an oil of the title compound (7.30 g).

[0100] Step 2: Preparation of 1-{2-[4-(2-Methoxyphenyl)-piperazin-1-yl]-ethyl}-1,5,6,7-tetrahydro-indol-4-one

[0101] A mixture of 1-(2-Chloroethyl)-4-(2-methoxyphenyl)piperazine (7.30 g) was dissolved in DMSO (30 mL), then slowly added to 1,5,6,7-tetrahydro-indol-4-one (30.1 mmol) which was pretreated with sodium hydroxide (31.6 mmol) in DMSO (15 mL) for 0.5 h with slight heat. The reaction was left under heat and was monitored by TLC (ethyl acetate: dichloromethane, 1:1). After completion (8 h), the reaction mixture was poured in ice water (300 mL) and extracted with dichloromethane twice, dried with sodium sulfate and solvent removed under vacuum. Flash chromatography (ethyl acetate: dichloromethane, 1:4) yielded an oil, NEO-377 (7.25 g).

EXAMPLE 8

Synthesis of 6-Chloro-9-{3-[4-(3-trifluoromethylphenyl)piperazin-1-yl]propyl}-9H-purine

[0102] Step 1: Preparation of 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine

[0103] Preparation of 1-(3-Chloropropyl)-4-(3-trifluoromethylphenyl)piperazine is prepared by the same method as disclosed in step 1 of Example 2 above.

[0104] Step 2: Preparation of 6-Chloro-9-{3-[4-(3-trifluoromethylphenyl)piperazin-1-yl]propyl}-9H-purine

[0105] Sodium hydride (60% in oil) (85 mg, 2.1 mmol, 1.8 eq.) was added to a 10 ml pear-shaped flask. The solid was rinsed twice with 2 mL hexane to remove oil, then potassium carbonate was added. 1,5,6,7-Tetrahydroindol-4-one (186.7 mg, 1.381 mmol, 1.159 eq.) was added slowly, with stirring and hydrogen evolved. The walls of the flask were washed with an additional potassium carbonate. 1-(2-Chloro-propyl)-4-(3-trifluoromethyl-phenyl)-piperazine (349.0 mg, 1.192 mmol, 1.000 eq) was added as a solution in 2 mL DMF, and the mixture was stirred under nitrogen at 25°C for 8 hours. The resulting mixture was acidified with 1 N HCl to pH 6, and extracted with dichloromethane. The organic layer was washed four times with 25 mL water, dried over sodium sulfate and concentrated in vacuum to an oil which was purified by column chromatography using 5% methanol in dichloromethane as eluant resulting in the title compound as an oil. The oil was dissolved in 5 mL of 50% dichloromethane in hexanes. A solution of 4N HCl in dioxane (200 µL) was added and the mixture stirred for 30 minutes followed by vacuum filtration of the suspension. A white powder of the product HCl salt was recovered.

EXAMPLE 9

Synthesis of 9-{3-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]propyl}-1,9-dihydropurin-6-one

[0106] 6-Chloro-9-{3-[4-(3-trifluoromethylphenyl)piperazin-1-yl]propyl}-9*H*-purine (200 mg) was dissolved in 3mL of trifluoroacetic acid(TFA) and 1 mL of water and stirred for 16 hours at 25 C. The TFA was evaporated and 10 mL of 2N NaOH was added. The solution was extracted twice with 20 mL of dichloromethane. The organic layer was separated, dried with sodium sulfate and concentrated to give the title compound as an oil. The oil was dissolved in 5 mL of 50% dichloromethane in hexanes. A solution of 4N HCl in dioxane (200 uL) was added and the mixture stirred for 30 minutes followed by vacuum filtration of the suspension. A white powder of the product HCl salt was recovered.

EXAMPLE 10

Synthesis of 1-{4-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]butyl}-1,5,6,7-tetrahydroindol-4-one (NEO-392)

[0107] Step 1: Synthesis of 1-(4-Chlorobutyl)-1,5,6,7-tetrahydroindol-4-one:

To a solution of 1,5,6,7-tetrahydroindol-4-one (10.0 g, 74.0 mmol) in acetone (300 mL) was added powdered sodium hydroxide (3.26 g, 81.4 mmol) and the mixture

stirred at ambient temperature for 0.25 h. 1-Bromo-4-chlorobutane (9.38 mL, 81.4 mmol) was then added and the resulting mixture stirred at ambient temperature for 7 h after which time TLC (ethyl acetate:dichloromethane 1:1) showed complete reaction. The reaction was gravity filtered to remove salts, and the filtrate concentrated to dryness under vacuum. The resulting residue was dissolved in dichloromethane (200 mL) and gravity filtered again to remove more salts. The filtrate was then washed with water, dried with sodium sulfate, filtered and the solvent removed under vacuum to yield an oil. Flash chromatography using 6 in. of silica gel in a 5.5 cm column eluting with 1:1 followed by 2:1 ethyl acetate:hexane on half of the residue yielded 9.0 g of an oil which contained ~6.0 g of pure product (72%) and ~3.0 g of acetone aldol condensation product (4-hydroxy-4-methyl-2-pentanone). The oil was taken to the next step without further purification.

Step 2: Synthesis of 1-{4-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]butyl}-1,5,6,7-tetrahydroindol-4-one

A mixture of 1-(4-Chlorobutyl)-1,5,6,7-tetrahydroindol-4-one (6.0 g, 26.6 mmol, as a mixture with 3.0 g of 4-hydroxy-4-methyl-2-pentanone) and sodium iodide (4.38 g, 29.2 mmol) in acetonitrile (100 mL) was heated at reflux for 6 h. (3-Trifluoromethylphenyl) piperazine (5.81 g, 25.2 mmol) and potassium carbonate (3.67 g, 26.6 mmol) was then added and reflux continued for 16 h. TLC (ethyl acetate:dichloromethane 1:1) showed complete reaction. The reaction was poured into ice cold water (400 mL) and stirred for 0.5 h. An oil separated out and was isolated from the mixture. The oil was dissolved with dichloromethane (150 mL), washed with water and brine, then dried with sodium sulfate, filtered and the solvent removed under vacuum to yield the title compound as an oil (9.7 g, 91.5%).

Preparation of Oxalate salt of 1-{4-[4-(3-Trifluoromethylphenyl)piperazin-1-yl]butyl}-1,5,6,7-tetrahydroindol-4-one (NEO-392). Dissolved compound (4.2 g) in hot ethyl acetate (150 mL), filtered solution hot to remove undissolved solid, and added a solution of oxalic acid (1.08 g, 1.2 eq) in methanol (10 mL) with stirring. A white precipitate formed immediately and the mixture was stirred for 0.5 h to room temperature. Vacuum filtration and washing with ethyl acetate afforded an off-white powder upon drying (5.0 g, 98%). HPLC Purity: 98.9%.

III. METHODS OF USING TETRAHYDROINDOLONE, PURINE OR DERIVATIVES THEREOF LINKED TO ARYLPIPERAZINES ACCORDING TO THE PRESENT INVENTION

[0108] The present invention includes methods of treating psychiatric and neurological disorders while enhancing cognitive function. This unexpected result is achieved by administering a compound having multiple pharmacophore components contained within the composite molecule. The multiple pharmacophores include tetrahydroindolone or purine derivatives linked to arylpiperazine structures. The term "pharmacophore," as used herein refers to a structural component of a molecule that causes a pharmacological response. In particular, tetrahydroindolone or purine derivatives incorporating arylpiperazine groups can be used as anti-psychotic compounds and administered to treat psychiatric disorders such as depression, anxiety, schizophrenia, schizoaffective disorders, bipolar disorders, sexual dysfunction, mood swings, sleep disorders, anorexia, bulimia, manic depression, obsessive compulsive disorders, delusional post-partum depression, post-partum psychosis, pre-menstrual syndrome, drug abuse associated psychoses and combinations thereof; neuroregenerative disorders with cognitive deterioration such as Parkinson's disease, Huntington's disease, Alzheimer's disease, and dementia associated with aging; pain associated with a wide range of conditions including drug induced neuropathy; and genetic disorders.

[0109] The therapeutic effect of the composite compound of the present invention is achieved by the compound's ability to affect multiple neuroreceptors. That is, by providing multiple pharmacophores having different receptor activities, the present invention is able to achieve therapeutic benefits unattainable by current therapies. For example, the arylpiperazine derivatives described as Moiety B of the composite compounds affect serotonin (5-HT) and dopamine receptors. In particular, the arylpiperazine derivatives contained in the composite compound have 5-HT_{1A} receptor agonist activity and can have activity at other receptors including, but not limited to, 5-HT₁ (A-F), 5-HT₂ (A-C), 5-HT₃ (1-7), 5-HT₄ C, 5-HT₅ (A-B) and dopaminergic receptors including, but not limited to D₂, D₃, and D₄. With all of the activities defined by this pharmacological profile, the composite compound will possess antipsychotic properties.

[0110] Additionally, the tetrahydroindolone or purine derivative component comprising moiety A of the composite compounds believed to be a pharmacophore

predicted to have GABA activity. As those skilled in the art will appreciate, GABA receptors are highly localized in the hippocampal region of the brain which is associated with memory. Generally recognized GABA receptors include, but are not limited to GABA A α (1-6), GABA A β (1-3), GABA A γ (1-3), GABA A δ , GABA A π , GABA A θ , GABA A ρ (1-3), GABA B1 (a-c), GABA B2, and GABA C. Accordingly, it is assumed that pharmacophores having GABA activity will enhance cognitive function. Thus, by providing a composite compound comprising tetrahydroindolone or purine derivatives linked with arylpiperazines, psychiatric disorders can be treated without causing a decrease in cognitive function.

[0111] In another embodiment of the present invention the pharmaceutical compositions disclosed herein selectively modulates 5HT1 and GABA and/or dopamine receptors. By “selectively modulates” the present inventors mean a compound that interacts preferentially with a receptor causing increases or decreases in related neurological functions a compared with other receptors.

[0112] Given the ability of the composite compounds of the present invention to affect activity at multiple receptors it is contemplated that other beneficial effects can be realized. For example, the composite compounds can be useful for motion sickness or drug-induced nausea. Moreover, because the composite compounds of the present invention has 5-HT1A receptor agonist activity the composite compounds can be useful in treating neurodegenerative disorders. As previous stated the novel composite compounds of the present invention can be useful as pharmaceutical compositions treating a number of psychiatric and neurological diseases. Psychiatric disorders include, but are not limited to, anxiety, depression, psychosis, schizoaffective, bipolar disorders, sexual dysfunction, mood swings and related cognitive defects including attention deficient disorders. Furthermore, anxiety, depression and psychosis induced by drugs, diseases and injury can also be effectively treated using the pharmaceutical compositions of the present invention.

[0113] Emotional, mood swings and cognitive disorders related to psychiatric disturbances that are expressed as sleep disorders, anorexia, bulimia, manic depression, obsessive compulsive disorders, and delusional disorders can be treated using the pharmaceutical compositions of the present invention. Moreover, the present inventors have also determined that other, more transient emotional disorders such as post-partum depression, post-partum psychosis, and pre-menstrual syndrome also benefit from the pharmaceutical compositions disclosed herein.

[0114] Other emotional disturbances that can be effectively treated include those related to substance abuse. For example, the pharmaceutical compositions of the present invention can be used prevent drug dependence or tolerance. Furthermore, the pharmaceutical compositions of the present invention can be useful in preventing or treating emotional and cognitive disturbances or psychoses associated with drug withdrawal or cessation.

[0115] Neurological diseases that can be effectively treated using the teachings of the present invention include disorders such as, but not limited to, neurosensory diseases and injury, Parkinson's disease and other movement disorders such as dystonia, Huntington's disease, Wilson's disease, inherited ataxias, Tourette syndrome cerebral palsy, encephalopathies and the related cognitive and emotional disorders associated therewith.

[0116] Pharmaceutical compositions made in accordance with the teachings of the present invention are also useful in treating a wide range of cognitive disorders; attention deficits and age related dementia in addition to those associated with the psychiatric disorders discussed above. For example, and not intended as a limitation, cognitive and attention deficit disorders associated with acquired immunodeficiency syndrome (AIDS), Down's syndrome, hyperactivity disorder, Alzheimer's disease and dementia, ischemic stroke, and cardiac bypass associated cognitive defects are also treatable using the pharmaceutical compositions described herein.

[0117] In another embodiment of the present invention the pharmaceutical compositions disclosed and claimed herein can also useful as treatments for pain associated with a wide range of conditions. For example, acute pain associated with injury or trauma and chronic pain lasting for periods in excess of six months are both treated using the pharmaceutical compositions of the present invention. Pain can be divided into two broad categories: nociceptive and neuropathic (non-nociceptive). These types of pain differ in their causes, symptoms, and responses to analgesics. Nociceptive or somatic pain results from direct stimulation of nociceptive, intact (uninjured) afferent nerve endings. Descriptors for this type of pain are usually "dull," "sharp," and/or "aching," or a combination of these, and the intensity of the pain varies from mild to severe. In general, somatic pain can be well controlled if the cause of the stimulation can be removed or otherwise treated (surgery, radiation therapy, chemotherapy, etc.), or treated with analgesics.

[0118] Neuropathic pain, on the other hand, is caused by nervous system dysfunction rather than stimulation of intact afferent nerve endings. It is characterized by burning, shooting, and tingling pain, associated with allodynia, hyperpathia, paresthesias and dysesthesias. Moreover neuropathic pain is often non-responsive to analgesics as in the case of opioid resistant neuropathic pain.

[0119] Although it is difficult to generalize when categorizing pain types and sources, the following general, non-limiting pain categories treatable using the pharmaceutical compositions of the present invention include both nociceptive and neuropathic pain. For example, nociceptive pain includes, without limitation, pain associated with arthritis, AIDS, chronic back pain, visceral organs, gastroesophageal reflux, peptic ulcers, infectious gastritis, inflammatory bowel disorders, migraine headache, cluster headache, tension headache, fibromyalgia, nerve root compression such as sciatica, trigeminal neuralgia, central pain, bone injury pain, pain during labor and delivery, post-operative pain, dental pain, genito-urinary tract pain, burn pain, angina pain, muscle strain, alcoholism, herpetic neuralgia, phantom limb pain, and dysmenorrhea.

[0120] It has also been determined by the present inventors that the pharmaceutical compositions disclosed herein also possess neuroregeneration/neuroprotective effects in addition to antiemetic properties. For example, the present invention can be used for the treatment or prevention of the pathology of peripheral neuropathy, neurodegenerative aspects of schizophrenia, psychosis, depression, anxiety, mood swings, Down's syndrome, stroke as well as other neurodegenerative diseases that can be genetic, spontaneous or iatrogenic including, but not limited to spinal cord injury amyotrophic lateral sclerosis, perinatal hypoxia, ocular damage and retinopathy, ocular nerve degeneration, hearing loss, restless leg syndrome and Tourette's syndrome.

[0121] The composite compounds of the present invention can be used to treat peripheral neuropathies and the neuropathic pain associated with peripheral neuropathies. Peripheral neuropathies can be genetic, or induced by diseases and/or drugs. Examples of disease associated with peripheral neuropathies include, but are not limited to acromegaly, hypothyroidism, AIDS, leprosy, Lyme disease, systemic lupus erythematosus, rheumatoid arthritis, Sjögren's Syndrome, periarteritis nodosa, Wegener's granulomatosis, cranial arteritis, sarcoidosis, diabetes, vitamin B12 deficiency, cancer and alcoholism. Examples of drug therapies associated with peripheral neuropathies include, but are not limited to oncolytic drugs such as a vinca

alkaloid, cisplatin, paclitaxel, suramin, altretamine, carboplatin, chlorambucil, cytarabine, dacarbazine, docetaxel, etoposide, fludarabine, ifosfamide with mesna, tamoxifen, teniposide, or thioguanine. Methods according to the present invention are particularly useful in treating drug-induced peripheral neuropathy arising from the administration of vinca alkaloids, taxanes, or platinum derivatives.

[0122] Depending upon the particular needs of the individual subject involved, the composite compounds of the present invention can be administered in various doses to provide effective treatment concentrations based upon the teachings of the present invention. Factors such as the activity of the selected composite compounds, the physiological characteristics of the subject, the extent or nature of the subject's disease or condition, and the method of administration will determine what constitutes an effective amount of the selected composite compounds. Generally, initial doses will be modified to determine the optimum dosage for treatment of the particular mammalian subject. The composite compounds can be administered using a number of different routes including orally, topically, transdermally, intraperitoneal injection, or intravenous injection directly into the bloodstream. Of course, effective amounts of the composite compounds can also be administered through injection into the cerebrospinal fluid or infusion directly into the brain, if desired.

[0123] An effective amount of any embodiment of the present invention is determined using methods known to pharmacologists and clinicians having ordinary skill in the art. For example, a pain relieving effective amount can be determined subjectively by administering increasing amounts of the pharmaceutical compositions of the present invention until such time the patient being treated reports diminishment in pain sensations. Blood levels of the drug can be determined using routine biological and chemical assays and these blood levels can be matched to the route of administration. The blood level and route of administration giving the most desirable level of pain relief can then be used to establish an "effective amount" of the pharmaceutical composition for treating the pain under study. This same method of titrating a pharmaceutical composition in parallel with administration route can be used to ascertain an "effective amount" of the pharmaceutical compositions of the present invention for treating any and all psychiatric or neurological disorders described herein.

[0124] Exemplary dosages in accordance with the teachings of the present invention for these composite compounds range from 0.0001 mg/kg to 60 mg/kg, though

alternative dosages are contemplated as being within the scope of the present invention. Suitable dosages can be chosen by the treating physician by taking into account such factors as the size, weight, age, and sex of the patient, the physiological state of the patient, the severity of the condition for which the composite compound is being administered, the response to treatment, the type and quantity of other medications being given to the patient that might interact with the composite compound, either potentiating it or inhibiting it, and other pharmacokinetic considerations such as liver and kidney function.

EXAMPLES

[0125] The following non-limiting examples demonstrate the clinical utility of the present invention. Specifically, Example 11 demonstrates the present invention's bioavailability in epithelial cells compared to two United States Food and Drug administration approved antipsychotic drugs (clozapine and risperidone). Example 12 demonstrates the anti-psychotic efficacy of the present invention in mice using pre-pulse inhibition testing. In Example 13 the anti-psychotic efficacy of the present invention is demonstrated in mice using conditioned avoidance testing (CAR).

EXAMPLE 11

Membrane Permeability

[0126] Caco-2 cells were used to evaluate the membrane permeability of eight embodiments of the present invention. Monolayers were grown to confluence and the integrity checked by transepithelial electrical resistance (TEER) and permeability of Lucifer Yellow. The expression of P-glycoprotein was confirmed by measuring the permeability of digoxin alone and in the presence of verapamil. Permeability measurements were taken for each compound in the apical to basolateral (AP-BL) direction, and the basolateral to apical (BL-AP) direction. Results are presented in Table 1 below.

Table 1: Comparative Membrane Permeability (P_{app}) Across Caco-2 Cells

CODE	P_{app} (cm/s x 10^6) AP- to- BL	P_{app} (cm/s x 10^6) BL- to- AP	RATIO $\frac{P_{app}^{B-A}}{P_{app}^{A-B}}$	PERMEABIL- ITY	EFFLUX LIMITED	POTENTIAL TRANSPOR- TER MEDIATED
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Propranolol	17.3±2.3			High	No	
Atenolol	1.3±0.2			Low	No	
NEO-356	21.5±2.9	8.8±0.3	0.41	High	No	+
NEO-359	24.0±3.9	7.6±0.5	0.32	High	No	+
NEO-363	18.0±0.7	13.1±1.1	0.75	High	No	-
NEO-370	32.1±8.1	20.0±3.4	0.67	High	No	-
NEO-376	20.4±1.8	9.4±0.3	0.43	High	No	+
NEO-377	26.0±1.8	15.5±2.9	0.60	High	No	-
NEO-381	32.9±9.6	25.3±0.7	0.77	High	No	-
NEO-392	27.1±1.2	15.7±1.3	0.58	High	No	-
Clozapine	15.3±0.7	7.4±1.1	0.65	High	No	-
Risperidone	21.5±0.2	14.0±3.1	0.52	High	No	-

EXAMPLE 12

Pre-pulse Inhibition Testing

[0127] The non-competitive NMDA receptor antagonist phencyclidine (PCP) reduces pre-pulse inhibition (PPI) of the acoustic startle response in rodents. Identifying compounds that improve the PCP-induced deficits in pre-pulse inhibition may provide insight into therapeutic strategies for treating schizophrenia. For testing of PPI, male C-57 mice were assigned to five dose groups of eight animals per group, and vehicle or test compound were administered orally (PO) or subcutaneously (SC) 20 minutes prior to intraperitoneal (IP) administration of vehicle or PCP (5mg/kg). Ten minutes following PCP administration, the mice were placed into Hamilton-Kinder startle

chambers and evaluation of pre-pulse inhibition procedure was performed. Following a five-minute acclimatization period with background white noise (65db), mice were exposed to five different trial types. Trials were presented ten-time search in a quasi-random order, with randomized 5 to 25 second inter-trial intervals. Trials were: stimulus only trial (120db white noise, 50ms stimulus); two different prepulse+pulse trials in which a 20ms 5db, or 10db stimuli above a 65db background preceded the 120db pulse by 120ms; a 10dbprepulse without a 120db pulse; and a no stimulus trial, in which only the background noise was presented. Results are presented in Table 2 below.

Table 2 PCP Pre-pulse Inhibition

Compound	PCP Disruption of Pre-pulse Inhibition	PCP Disruption of Pre-pulse Inhibition – SC
NEO-356 (oxalate)	No reversal	5 pp ED50=2.31 mg/kg
NEO-359 (oxalate)	No reversal	5pp ED50=3.58mg/kg 10pp ED50=0.79mg/kg
NEO-363 (oxalate)	10pp ED50=3.10 mg/kg	5pp Ed=2.30 mg/kg 10pp ED50=2.30 mg/kg
NEO-370 (oxalate)	5pp & 10pp ED50=0.96 mg/kg	5pp ED50=0.79 mg/kg 10pp ED50=0.79 mg/kg
NEO-376 (oxalate)	10pp ED50=0.79 mg/kg	5pp ED50=0.79 mg/kg 10pp ED50=0.79 mg/kg
NEO-377 (oxalate)	10pp ED50=2.30 mg/kg	5pp ED50=2.31 mg/kg 10pp ED50=1.12 mg/kg
NEO-381 (oxalate)	No reversal	5pp ED50=0.79 mg/kg 10pp ED50=3.58 mg/kg
NEO-392 (oxalate)	10pp ED50=1.12 mg/kg	5pp ED50=0.23 mg/kg 10pp ED50=0.23 mg/kg
Clozapine		5pp ED50=2.87 mg/kg 10pp ED50=3.00 mg/kg
Risperidone		5pp ED50=0.036 mg/kg 10pp ED50=0.036 mg/kg
***PCP-Induced Psychosis (Disruption of Pre-pulse Inhibition): Reversal of disruption of pre-pulse inhibition produced by PCP is a clinical predictor of compounds with antipsychotic activity		
Note: estimated ED50 values		

EXAMPLE 13

Conditioned Avoidance Testing

[0128] Training: Consists of 20 trials with variable inter-trial intervals (Trained to 80% Avoidance Criteria). After a one-minute acclimation period, the house light and an acoustic 90dB tone (conditioned stimuli) are presented. A response (crossing to dark compartment) within 5 seconds ends the trial and trial is recorded as avoidance response (CAR). If the mouse does not respond within 5 seconds, foot shock (0.8 mA) is presented, and the response (moving to the dark chamber) during the shock was recorded as an escape response. To avoid shock, animals learn to move from the lighted side of the chamber to the dark side when the cue is presented (avoidance) or moved when the shock is administered (escape). Testing: Vehicle or test compounds are administered S.C. 20 minutes before the test session. (Checking for disruption of cognition and attention)

Table 3. Conditioned Avoidance Responding

Compound	Conditioned Avoidance Responding**
NEO-356	No effect
NEO-359	No effect on escape or avoidance Significant decrease in latency <i>latency ED50=100.00 mg/kg</i>
NEO-363	No effect
NEO-370	Significant increase in latency Significant reduction in avoidance <i>Latency ED50=3.25 mg/kg; avoidance ED50=6.20 mg/kg</i>
NEO-376	No effect on escape or avoidance significant increase in latency <i>Latency ED50= 17.49 mg/kg</i>
NEO-377	Significant increase in latency Significant reduction in avoidance <i>Latency ED50=3.0 mg/kg ; avoidance ED50=7.57 mg/kg</i>
NEO-381	No effect
NEO-392	Significant increase in latency Significant reduction in avoidance <i>Latency ED50=3.25 mg/kg; avoidance ED50=4.93 mg/kg</i>
Clozapine	Significant increase in latency Significant reduction in avoidance <i>Latency ED50=3.03 mg/kg ; avoidance ED50 = 6.30 mg/kg</i>

Risperidone	Significant increase in latency Significant reduction in avoidance <i>Latency ED50=0.26 mg/kg</i> <i>Avoidance ED50=0.19 mg/kg</i>
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In all behavioral experiments shown drug was administered SC to mice, unless otherwise noted.

****Condition Avoidance Responding (CAR, active avoidance):** disruption of avoidance (increased latency) without disruption of escape (extrapyramidal motor function) is a clinical predictor of compounds with antipsychotic activity

IV. PHARMACEUTICAL COMPOSITIONS

[0129] Another aspect of the present invention is a pharmaceutical composition that comprises: (1) an effective amount of a composite compound according to the present invention as described above; and (2) a pharmaceutically acceptable excipient.

[0130] A pharmaceutically acceptable excipient, including carriers, can be chosen from those generally known in the art including, but not limited to, inert solid diluents, aqueous solutions, or non-toxic organic solvents. If desired, these pharmaceutical formulations can also contain preservatives and stabilizing agents and the like, as well as minor amounts of excipients substances such as, but not limited to, a pharmaceutically acceptable excipient selected from the group consisting of wetting or emulsifying agents, pH buffering agents, human serum albumin, ion exchanger resins, antioxidants, preservatives, bacteriostatic agents, dextrose, sucrose, trehalose, maltose, alumina, lecithin, glycine, sorbic acid, propylene glycol, polyethylene glycol, protamine sulfate, sodium chloride, or potassium chloride, mineral oil, vegetable oils and combinations thereof. Those skilled in the art will appreciate that other carriers also can be used.

[0131] Liquid compositions can also contain liquid phase excipients either in addition to or to the exclusion of water. Examples of such additional liquid phases are glycerin, vegetable oils such as cottonseed oil, organic esters such as ethyl oleate, and water-oil emulsions.

[0132] Formulations suitable for parenteral administration, such as, for example, by intravenous, intramuscular, intradermal, and subcutaneous routes, include aqueous and non-aqueous isotonic sterile injection solutions. These can contain antioxidants, buffers, preservatives, bacteriostatic agents, and solutes that render the formulation isotonic with the blood of the particular recipient. Alternatively, these formulations can be aqueous or non-aqueous sterile suspensions that can include suspending agents,

thickening agents, solublizers, stabilizers, and preservatives. The pharmaceutical compositions of the present invention can be formulated for administration by intravenous infusion, oral, topical, intraperitoneal, intravesical, transdermal, intranasal, intrarectal, intravaginal, intramuscular, intradermal, subcutaneous and intrathecal routes.

[0133] Formulations of composite compound suitable for use in methods according to the present invention can be presented in unit-dose or multi-dose sealed containers, in physical forms such as ampules or vials. The compositions can be made into aerosol formations (i.e., they can be “nebulized”) to be administered via inhalation. Aerosol formulations can be placed into pressurized acceptable propellants, such as dichloromethane, propane, or nitrogen. Other suitable propellants are known in the art.

[0134] The terms “a” and “an” and “the” and similar referents used in the context of describing the invention (especially in the context of the following claims) are to be construed to cover both the singular and the plural, unless otherwise indicated herein or clearly contradicted by context. Recitation of value ranges herein are merely intended to serve as a shorthand method of referring individually to each separate value falling within the range. Unless otherwise indicated herein, each individual value is incorporated into the specification as if it were individually recited herein. All methods described herein can be performed in any suitable order unless otherwise indicated herein or otherwise clearly contradicted by context. The use of any and all examples, or exemplary language (e.g. “such as”) provided herein is intended merely to better illuminate the invention and does not pose a limitation on the scope of the invention otherwise claimed. No language in the specification should be construed as indicating any non-claimed element essential to the practice of the invention.

[0135] Groupings of alternative elements or embodiments of the invention disclosed herein are not to be construed as limitations. Each group member can be referred to and claimed individually or in any combination with other members of the group or other elements found herein. It is anticipated that one or more members of a group can be included in, or deleted from, a group for reasons of convenience and/or patentability. When any such inclusion or deletion occurs, the specification is herein deemed to contain the group as modified thus fulfilling the written description of all Markush groups used in the appended claims.

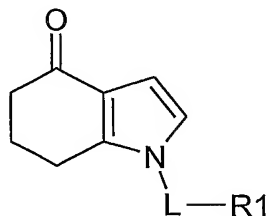
[0136] Preferred embodiments of this invention are described herein, including the best mode known to the inventors for carrying out the invention. Of course, variations on those preferred embodiments will become apparent to those of ordinary skill in the art upon reading the foregoing description. The inventor expects skilled artisans to employ such variations as appropriate, and the inventors intend for the invention to be practiced otherwise than specifically described herein. Accordingly, this invention includes all modifications and equivalents of the subject matter recited in the claims appended hereto as permitted by applicable law. Moreover, any combination of the above-described elements in all possible variations thereof is encompassed by the invention unless otherwise indicated herein or otherwise clearly contradicted by context.

[0137] Furthermore, numerous references have been made to patents and printed publications throughout this specification. Each of the above cited references and printed publications are herein individually incorporated by reference in their entirety.

[0138] In closing, it is to be understood that the embodiments of the invention disclosed herein are illustrative of the principles of the present invention. Other modifications that can be employed are within the scope of the invention. Thus, by way of example, but not of limitation, alternative configurations of the present invention can be utilized in accordance with the teachings herein. Accordingly, the present invention is not limited to that precisely as shown and described.

What is claimed is:

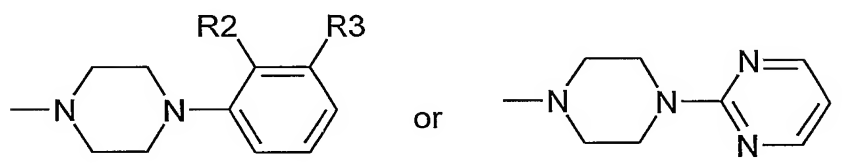
1. A pharmaceutical composition comprising:
an anti-psychotic compound having the structure of formula (XI);



(XI)

wherein L is $-(CH_2)_m-$ and m is an integer from 1 to 6;

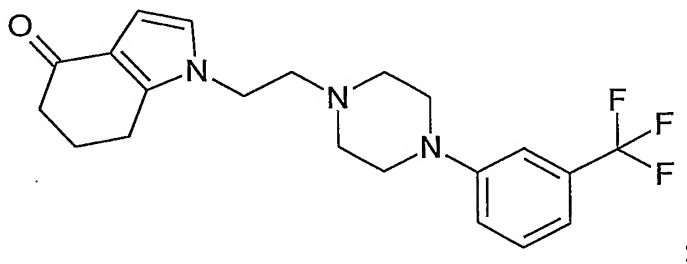
R1 is:



wherein R2 and R3 are the same or independently hydrogen, alkyl, hydroxy, methoxy, halo, alkoxy, trifluoromethyl, nitro, amino, aminocarbonyl, or aminosulfonyl;

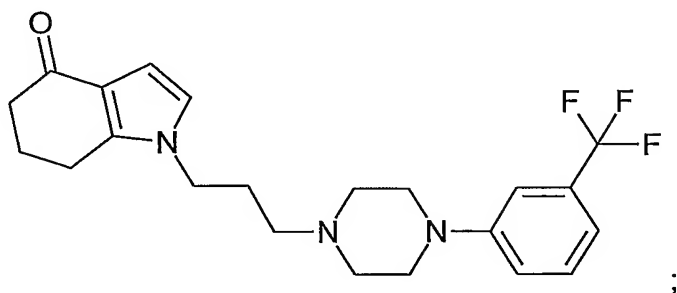
and pharmaceutically acceptable salts thereof.

2. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



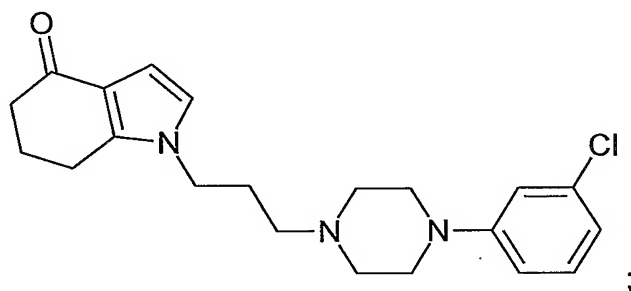
and pharmaceutically acceptable salts thereof.

3. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



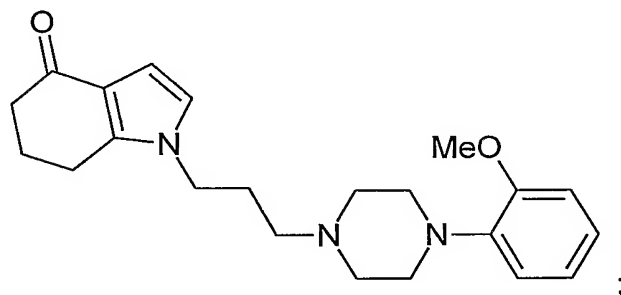
and pharmaceutically acceptable salts thereof.

4. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



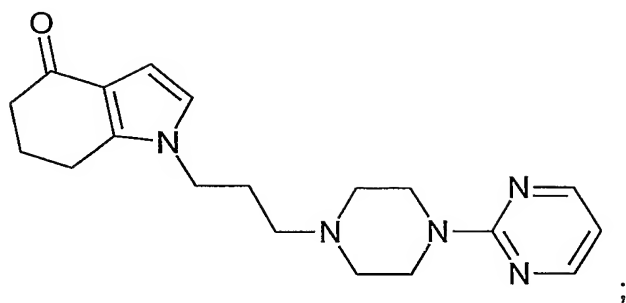
and pharmaceutically acceptable salts thereof.

5. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



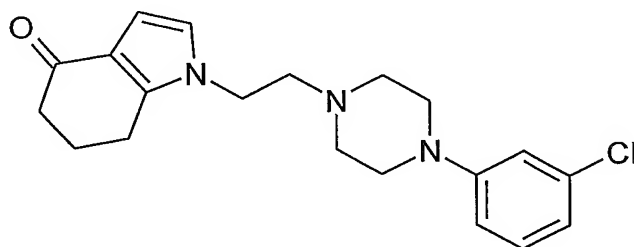
and pharmaceutically acceptable salts thereof.

6. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



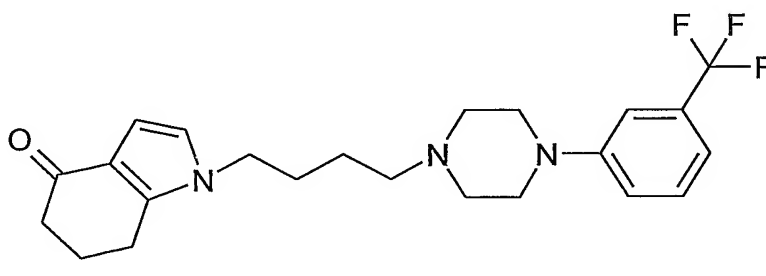
and pharmaceutically acceptable salts thereof.

7. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



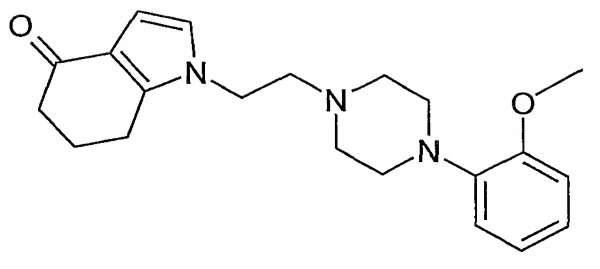
and pharmaceutically acceptable salts thereof.

8. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



and pharmaceutically acceptable salts thereof.

9. The pharmaceutical compositions according to claim 1 wherein said anti-psychotic compound comprises:



and pharmaceutically acceptable salts thereof.

10. The pharmaceutical compositions according to any one of claim 1 through 9 further comprising:

a pharmaceutically acceptable excipient selected from the group consisting of wetting or emulsifying agents, pH buffering agents, human serum albumin, ion exchanger resins, antioxidants, preservatives, bacteriostatic agents, dextrose, sucrose, trehalose, maltose, alumina, lecithin, glycine, sorbic acid, propylene glycol, polyethylene glycol, protamine sulfate, sodium chloride, or potassium chloride, mineral oil, vegetable oils and combinations thereof.

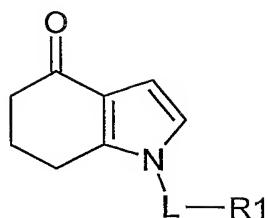
11. A pharmaceutical composition according to claim 10 wherein said pharmaceutical composition is formulated for administration by routes selected from the group consisting of intravenous infusion, oral, topical, intraperitoneal, intravesical, transdermal, intranasal, intrarectal, intravaginal, intramuscular, intradermal, subcutaneous and intrathecal

12. A method of treating a psychiatric disorder comprising:

administering a pharmaceutically effective amount of any compound according to claim 1 through 10.

13. The method according to claim 12 wherein said psychiatric disorder is selected from the group consisting of depression, anxiety, schizophrenia, schizoaffective disorders, bipolar disorders, sexual dysfunction, mood swings, sleep disorders, anorexia, bulimia, manic depression, obsessive compulsive disorders, delusional post-partum depression, post-partum psychosis, pre-menstrual syndrome, drug abuse associated psychoses and combinations thereof.

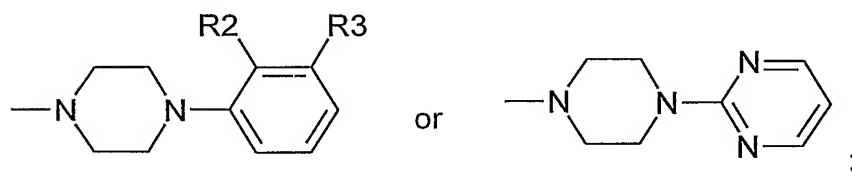
14. A composition that selectively modulates 5-HT₁, GABA and/or dopamine receptors comprising:



(XI)

wherein L is $-(CH_2)_m-$ and m is an integer from 1 to 6;

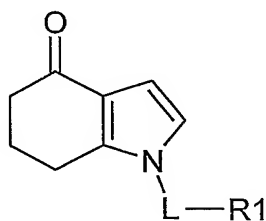
R1 is:



wherein R2 and R3 are the same or independently hydrogen, alkyl, hydroxy, methoxy, halo, alkoxy, trifluoromethyl, nitro, amino, aminocarbonyl, or aminosulfonyl;

and pharmaceutical salts thereof.

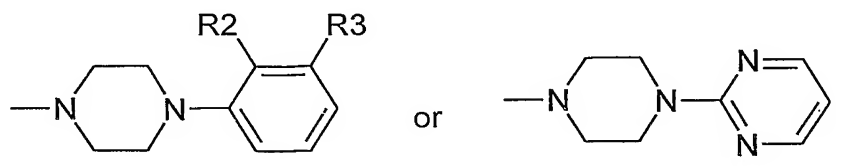
15. A compound useful for treating psychotic disorders comprising:



(XI)

wherein L is $-(CH_2)_m-$ and m is an integer from 1 to 6;

R1 is:



wherein R2 and R3 are the same or independently hydrogen, alkyl, hydroxy, methoxy, halo, alkoxy, trifluoromethyl, nitro, amino, aminocarbonyl, or aminosulfonyl;

and pharmaceutical acceptable salts thereof, providing, said compound does not impair cognitive functions.

INTERNATIONAL SEARCH REPORT

national Application No
PCT/US 02/24260

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61P25/00 A61P25/18 A61P25/22 A61P25/24 A61P15/00
A61P15/10 A61K31/496 A61K31/506 C07D403/12 C07D209/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61P A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, MEDLINE, EPO-Internal, BIOSIS, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 99 62899 A (BRYANT HELEN JANE ;MERCK SHARP & DOHME (GB); CURTIS NEIL ROY (GB);) 9 December 1999 (1999-12-09) the whole document	1-15
A	WO 93 12085 A (ASTRA AB) 24 June 1993 (1993-06-24) the whole document	1-15

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents : ...

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

31 October 2002

Date of mailing of the international search report

07/11/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
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Authorized officer

van der Kooij, M

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 02/24260

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 12 and 13 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

national Application No

PCT/US 02/24260

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
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			WO 9962899 A1	09-12-1999
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